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#### (54) Title: MODIFIED CELLULOSE ETHERS

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#### (57) Abstract

The invention relates to a cellulose ether that is hydrophobically modified with an alkyl or alkenylketene dimer of general formula (I), in which R<sub>1</sub> and R<sub>2</sub> are alkyl or alkenyl chain,

$$R_1 - CH = C - CH - R_2$$

| | | (1)

 $O - C = O$ 

in which R<sub>1</sub> and R<sub>2</sub> are alkyl or alkenyl chain, such as one having 5 to 22, preferably 13 to 20, most preferably 14 to 18 carbons. The product can be used in water-based products to control the rheological properties, for example. Objects of use include, for example, detergents, paper coating mixtures, paints, dispersing agents, and oil drilling muds.

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PCT/F199/00296

#### MODIFIED CELLULOSE ETHERS

### **Background of the Invention**

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The invention relates to hydrophobically modified cellulose ethers and to the use thereof.

Cellulose ethers, such as carboxymethyl cellulose, are employed in a wide variety of applications to treat the rheological properties of aqueous solutions in particular. Objects of use include, for example, foods, medicines, cosmetic preparations, detergents, agricultural chemicals, textiles, printing inks, paper coating agents, construction materials, adhesives, paints, ceramic materials, and additives of polymerisation.

Cellulose ethers can be modified by attaching different substituents to hydroxyl groups in cellulose. In this way, the solubility or the hydrophobicity of cellulose ether in particular can be affected.

There is a lot of literature concerning the hydrophobic modification of polysaccharides.

US Patent No. 4228277 describes the modification of non-ionic cellulose ethers, such as methyl cellulose, methyl hydroxypropyl cellulose, hydroxypropyl cellulose, and particularly hydroxyethyl cellulose, by using epoxyalkanes having a length of the alkyl group of 10 to 24 carbons. According to the patent, one advantage of the modification is that even small amounts of hydrophobically modified cellulose ether make the viscosity of an aqueous solution increase so that the dosage of the cellulose ether can be decreased. The cellulose ether thus modified can be used in paints. The modification is carried out by using a cellulose ether as a starting material, and the reaction times are long, 2 to 5 hours. However, it is uncomfortable to use epoxy compounds because they are detrimental to health.

In FI Patent No. 95138, carboxymethyl hydroxyethyl cellulose is modified by an alkyl group having 8 to 25 carbons in an amount of 0.1 to 4 % by weight. This product is also applied to paints. The alkyl reagent can be halide, halohydride, or epoxide, all of which are hazardous to environment and health. Hydroxyethylation, hydrophobic modification, and carboxymethylation are carried out in sequence, therefore the reaction time is long, more than 4 hours.

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US Patent No. 5302196 describes cellulose ethers which are modified by a fluorine-containing alkyl group having 3 to 24 carbon atoms, the fluorine content being 0.05 to 5% by weight. The cellulose ether is especially hydroxyethyl cellulose, hydroxyethyl carboxymethyl cellulose or methyl hydroxyethyl cellulose. The fluorine-containing alkyl reagent is preferably epoxide, bromide or perfluoroolefin. This product is also developed for paints. The reaction is long, the modification requires a reaction time of 6 hours.

In EP Patent Application No. 384167, cellulose ether, especially hydroxyethyl cellulose, is modified with an aromatic alkyl group having at least 10 carbons, the content being 0.001 to 0.1 moles per one mole of a glucose repeat unit of the cellulose ether. The alkyl reagent can be, for example, halide, oxirane, acid, (thio)isocyanate or halohydrin. The product can be used in latex paints. This manufacturing process also has a long reaction time, 6 or up to 12 hours.

# General Description of the Invention

15 According to the independent Claim 1, hydrophobically modified cellulose ethers have been invented. The dependent Claims disclose some preferred embodiments of the invention.

This invention introduces a new type of hydrophobically modified cellulose ether in which the modifying agent is alkylketene dimer. The cellulose ether is preferably carboxymethyl cellulose or sodium carboxymethyl cellulose (CMC, NaCMC).

Cellulose ether modified with alkylketene dimer is easy to manufacture. The process is safe, simple, and quick. The modification can also improve the dispersability of cellulose ether in water.

The product is suitable to be used in any water-based application. The hydrophobic group renders the substance properties that are advantageous to, for example, detergents, paper coating mixtures, paints, dispersing agents, and oil drilling muds.

# **Detailed Description of the Invention**

According to the invention, alkylketene dimer (AKD) is used to modify the cellulose ether. The general formula of AKD is 15

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$$R_1 - CH = C - CH - R_2$$
  
| | 0  
 $O - C = O$ 

5 in which R<sub>1</sub> and R<sub>2</sub> are alkyl or alkenyl chains having 5 to 22, preferably 13 to 20 carbons. R<sub>1</sub> or R<sub>2</sub> can also be substituted or they can contain heteroatoms in a chain. R<sub>1</sub> or R<sub>2</sub> can in particular have a straight chain and contain 14 to 18 carbons.

AKD reacts with the hydroxyl group of cellulose or cellulose ether, forming the following formula:

Elevated temperatures, such as 30-120°C, preferably 50-90°C, and most preferably 60-85°C, enhance the reaction, so that at the elevated temperatures AKD is melted first. High temperature and alkaline pH improve the reactivity of AKD with cellulose ether. AKD can react with the substituent, for example carboxymethyl group in cellulose ether. AKD can also react with other compounds which contain an OH group. These include water, alcohol, and sodium glycolate which are generated in the preparation of CMC. The thus generated compounds also have an effect on the hydrophobicity of the product.

The reacted AKD concentration in the cellulose ether can be analysed by using gas chromatography. The unreacted AKD or AKD reacted with other OH containing compounds are extracted with a suitable solvent. The sample is hydrolysed causing the cleavage of alkyl ketene group. The reacted AKD is extracted with a suitable solvent and analysed with gas chromatograph.

The AKD can be in a solid state or dispersed in water or other solvents.

The content of AKD in the cellulose ether can be 0.001-10% by weight. The content is preferably 0.01-2% by weight.

The molecular weight (Mw) of the cellulose ether can be 10 000 - 1 000 000, preferably 20 000 - 700 000. The cellulose ether can be, for example, alkyl, hydroxyalkyl or carboxyalkyl substituted or a mixed ether of these.

WO 99/61479 PCT/FI99/00296

According to the invention, cellulose ether that is modified by AKD can be used in any water-based application. It can be used, for example, in paper or board coating mixtures, in the wet end in paper or board manufacturing, paints, construction materials, adhesives, oil drilling mud, detergents, cosmetic products, and as a dispersion agent.

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The cellulose ether is preferably carboxymethyl cellulose (CMC). Its degree of substitution (DS, i.e., the average number of substituted hydroxyl groups in a glucose ring) can be, for example, 0.2-2.0, preferably 0.4-1.5, most preferably 0.4-1.2. Its molecular weight is preferably 40 000 - 500 000. It can be applied to all the above-mentioned objects.

The cellulose ether can also be, e.g., hydroxyethyl cellulose (HEC). Its molecular weight can be, for example, 90 000 - 1 300 000 and the molar substitution (MS) 1.5-4. HEC can be used in coating, construction materials, and cosmetic preparations in particular.

15 The cellulose ether can also be, e.g., hydroxypropyl cellulose (HPC). Its molecular weight can be, for example, 80 000 - 1 150 000 and the molar substitution (MS) 1.5-4. HPC can be used in foods, medical preparations, and coating in particular.

There is a lot of literature about the manufacture of cellulose ethers. Generally, cellulose ethers are prepared by mixing wood-based or cotton wool-based raw material with a reaction medium, such as alcohol or acetone, and by mercerising it with an alkalic substance, such as sodium hydroxide, to activate the cellulose. An etherifying chemical is added and it is allowed to react. The complete product is neutralised. The viscosity can be reduced if needed. By-products, such as salts generated in the reaction, are washed out with alcohol, for example, when purified cellulose ether is manufactured, but they can also be left in the product or only remove partly (technical or crude cellulose ether). The solvent used is separated and the product is dried. The particle size and the bulk density can be adjusted by grinding the product into powder or by granulating it. The product can also be screened.

Cellulose ether can be modified with AKD by various methods. AKD can be added to the cellulose ether at any stage of the production, for example, in mercerisation, etherification, or drying. AKD can be added in a solid form or as a dispersion. High temperature and alkaline pH improve the reactivity of AKD with cellulose ether.

AKD can also be added to the cellulose ether as such. The cellulose ether can be coated, for example, with solid AKD or an AKD dispersion. In that case, a suitable

WO 99/61479

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amount of AKD is sprayed onto the surface of the cellulose ether or mixed with it. The product is treated with high temperature to initiate the reaction.

The manufacture of the hydrophobically modified cellulose ether according to the invention is simple and quick. Neither complex solvents nor long reaction times are needed. In addition, AKD is a safe chemical. It is neither combustible nor explosive; therefore, it neither causes a danger in a process nor is it dangerous to the environment. AKD is not hazardous or irritating and it does not cause allergies; therefore, it does not present any danger to the process personnel or the end-user.

The hydrophobically modified cellulose ether is especially useful in detergent compositions. The cellulose ether is preferably carboxymethyl cellulose (CMC). Its degree of substitution is most preferably 0.4-0.6. The compositions generally comprise about 0,1-5% by weight of cellulosic based fabric treatment components, preferably 0,5-4%, most preferably 0,75-3%. The detergent composition can be in liquid, paste or granular form. The compositions comprise about 1-80% by weight, preferably 5-50% of detersive surfactants which can be of the anionic, nonionic, zwitterionic, ampholytic or cationic type. The compositions may also comprise about 0,1-80% by weight of a detergent builder, preferably 1-10% in liquid form and 1-50% in granular form. Detergent builders can comprise, for example, phosphate salts as well as various organic and inorganic nonphosphorus builders. In addition to the surfactants, builders and cellulosic based materials, the detergent compositions can also include for example enzymes and enzyme stabilizing agents, suds boosters or suds suppressers, anti-tarnish and anticorrosion agents, soil suspending agents, soil release agents, germicides, pH adjusting agents, non-builder alkalinity sources, chelating agents, organic and inorganic fillers, solvents, hydrotropes, optical brighteners, dyes and perfumes.

Granular detergent compositions are generally made by combining base ingredients as a slurry and spray drying the resulting slurry to a low level of residual moisture. The remaining ingredients can be admixed as dry or in case of liquid form, sprayed onto the resulting granules. Liquid detergent compositions can be prepared by admixing the ingredients in any desired order.

Laundering of fabrics and textiles in washing solutions containing hydrophobically modified cellulose ether, followed by rinsing and drying, imparts fabric appearance benefits to the fabric and textile articles so treated. Such benefits can include improved overall appearance, pill/fuzz reduction, antifading, improved abrasion resistance, and/or enhanced softness.

### **Examples**

All examples employed alkylketene dimer, the length of the alkyl group being 16 to 18 carbons (Raisio Chemicals).

### Example 1

100g of CMC (Metsa Speciality Chemicals) having an average molecular weight of 180 000 and a degree of substitution of 0.65 was put into a steel container. 1g of alkylketene dimer dispersion having an AKD content of 10% by weight, diluted with 50g of water, was sprayed on CMC, agitating the powder at the same time. The product was dried at 80°C to a moisture content of less than 8%. The AKD content of the end product was 0.1% by weight.

## Examples 2-6

By using the procedure described above, the following hydrophobically modified CMC samples (examples 2-6) were prepared:

Table 1

Example	Molecular weight of CMC	Degree of substi- tution of CMC	AKD content in CMC, % by weight
2	60 000	0.72	0.05
3	300 000	0.80	0.05
4	220 000	0.87	0.1
5	305 000	1.15	1.0
6	80 000	0.75	0.01

15 Sample 5 contained 0.02% by weight of AKD which had reacted with CMC.

# Example 7

100g of CMC having an average molecular weight of 40 000 and a degree of substitution of 0.79 was put into a steel cylinder. 5g of solid alkylketene dimer was added. The powder was agitated in an oven at 60°C for 120 minutes and at 105°C for 15 min. The AKD content of the end product was 5.0% by weight.

# Example 8

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Hydrophobically modified CMC having an average molecular weight of 400 000 and a degree of substitution of 0.86, and an AKD content of 10.0% by weight, was prepared according to the procedure described in example 7.

### Example 9

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47g of wood pulp (Metsä-Botnia), 175g of ethanol, and 20g of water were mixed together in a closed glass flask which was provided with a mixer and a nitrogen atmosphere as an option. The flask was placed in a water bath. 18g of sodium hydroxide, diluted with 18g of water, was added and the cellulose was mercerised for 30 minutes at 20°C. 22g of monochloracetic acid, diluted with 6g of water, and 1.25g of alkylketene dimer dispersion having an AKD content of 20% by weight were added and the temperature was raised in 55 minutes to 65°C. The cellulose was etherified for 60 minutes at 65°C. Ethanol was recovered and the product was dried at 80°C to a moisture content of less than 8%. The AKD content of the end product was 0.3% by weight, the average molecular weight 150 000, and the DS of the carboxymethyl group 0.57. The sample contained 0.02% by weight of AKD reacted with cellulose.

#### Examples 10-15

Samples 10-15 were prepared according to the procedure described in example 9. In example 10 and 15 the product was neutralised after etherification with hydrochloric acid. In example 10 AKD was added after neutralisation and allowed to react 5 minutes at 70 °C.

Table 2

Example	Molecular weight of CMC	Degree of substi- tution of CMC	AKD content in CMC, % by weight
10	180 000	0,46	0,6
11	120 000	0,48	0,6
12	140 000	0,52	0,06
13	150 000	0,55	1,1
14	140 000	0,56	2,7
15	270 000	0,52	0,06

20 Sample 12 contained <0.01% by weight of AKD which had reacted with CMC.

# Example 16

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100g of wood pulp (Metsä-Botnia), 1270g of isopropanol, and 162g of water were mixed together in a closed glass flask which was provided with a mixer and a nitrogen atmosphere. The flask was placed in a water bath. 40g of sodium hydroxide, diluted with 40g of water, was added and the cellulose was mercerised for 90 minutes at 20°C. 44g of monochloracetic acid, diluted with 11g of water, and 1g of al-

kylketene dimer dispersion having an AKD content of 10% by weight were added, and the temperature was raised in 45 minutes to 70°C. The cellulose was etherified for 90 minutes at 70°C. CMC was neutralised with hydrochloric acid. The byproducts of the reaction, sodium chloride and sodium glycolate, were washed out with 75% methanol, and the product was dried at 80°C to a moisture content of less than 8%. The AKD content of the end product was 0.08%, the average molecular weight 270 000, and the DS of the carboxymethyl group 0.59.

### Examples 17-19

Samples 17-19 were prepared according to the procedure described in example 16.

In example 19 AKD was added during mercerisation.

Table 3

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Example	Raw material	Molecular weight of CMC	Degree of substitution of CMC	AKD content in CMC, % by weight
17	wood pulp (Metsä-Botnia)	60 000	0,77	0,07
18	wood pulp (Borregaard)	240 000	0,89	0,07
19	cotton linter (Buckeye, Temming)	380 000	0,80	0,7

#### Example 20

A granular detergent composition in accordance with the invention has the following basic formula:

# Table 4

Component	Wt. %
C <sub>12</sub> Linear alkyl benzene sulfonate	9.31
C <sub>14-15</sub> alkyl ether (0.35 EO) sulfate	12.74
Zeolite Builder	27.79
Sodium Carbonate	27.31
PEG 4000	1.60
Dispersant	2.26
C <sub>12-13</sub> Alcohol Ethoxylate (9 EO)	1.5
Sodium Perborate	1.03
Soil Release Polymer	0.41
Enzymes	0.59
Hydrophobically modified cellulose ether	3.0
Perfume, Brightener, Suds Suppressor,	
Other Minors, Moisture, Sulfate	Balance
	100%

#### **CLAIMS**

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1. A hydrophobically modified cellulose ether, characterised in that the hydrophobic modification has been carried out by using an alkyl or alkenylketene dimer the general formula of which is

in which R<sub>1</sub> and R<sub>2</sub> are alkyl or alkenyl chains, such as one having 5 to 22, prefera-10 bly 13 to 20, most preferably 14 to 18 carbons.

2. A modified cellulose ether according to Claim 1, wherein the cellulose ether has the following formula:

O 
$$R_2$$
 O  $\parallel \ \mid \ \parallel$   $R_1$  -  $CH_2$  -  $C$  -  $CH$  -  $C$  -  $O$  - cellulose (ether)

in which R<sub>1</sub> and R<sub>2</sub> are as above.

- 3. A modified cellulose ether according to Claim 1 or 2, wherein the molecular weight of the cellulose ether is 10 000 1 000 000, preferably 20 000 700 000, most preferably 40 000 500 000.
  - 4. A modified cellulose ether according to any of Claims 1 to 3, wherein the content of the alkyl or alkenylketene dimer in the cellulose ether is 0.001-10% by weight, preferably 0.01-2% by weight.
- 5. A modified cellulose ether according to any of Claims 1 to 4, wherein the cellulose ether is carboxymethyl cellulose.
  - 6. A modified cellulose ether according to Claim 5, wherein the degree of substitution of the carboxymethyl cellulose is 0.2-2, preferably 0.4-1.5, and most preferably 0.4-1.2.
- 7. A composition intended to be mixed with water, or an aqueous composition, characterized in that the composition contains a hydrophobically modified cellulose ether according to any of Claims 1 to 6.
  - 8. A detergent composition according to Claim 7.

WO 99/61479 PCT/F199/00296

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- 9. A detergent composition according to Claim 8, wherein the composition contains hydrophobically modified carboxy methyl cellulose.
- 10. A detergent composition according to Claim 9, wherein the degree of substitution of carboxymethyl cellulose is 0.4-0.6.

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C DOCUM	ENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the rel	evant passages	Relevant to claim No.
X	WO 99 61479 A (METSA SPEC CHEM ON 2 December 1999 (1999-12-02)	ĺ	1-10
	page 4, line 11 - line 17; claims page 5, line 9 - line 35	,	
X	WO 00 65015 A (PROCTER & GAMBLE) 2 November 2000 (2000-11-02)		1-10
	page 2, line 16 - line 25; claims; examples page 13, line 16 - line 22		
	page 18, line 8 - line 26		
<b>X</b> (	WO 98 29528 A (PROCTER & GAMBLE) 9 July 1998 (1998-07-09) page 3, line 10 -page 8, line 16;	· claims	1–10
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#### INTERNATIONAL SEARCH REPORT

International application No.

PCT/FI 99/00296

# A. CLASSIFICATION OF SUBJECT MATTER According to International Patent Classification (IPC) or to both national classification and IPC IPC6: C08B 13/00 B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC6: C08B, C08L Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched SE,DK,FI,NO classes as above Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) CA, WPI, PAJ C. DOCUMENTS CONSIDERED TO BE RELEVANT Relevant to claim No. Citation of document, with indication, where appropriate, of the relevant passages Category\* 1-10 US 2627477 A (WILLIAM F. DOWNEY), 3 February 1953 X (03.02.53)US 3931069 A (LUNDIN), 6 January 1976 (06.01.76) 1-10 X See patent family annex. Further documents are listed in the continuation of Box C. later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive "F" erlier document but published on or after the international filing date document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) step when the document is taken alone "Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination document referring to an oral disclosure, use, exhibition or other heing obvious to a person skilled in the art document published prior to the international filing date but later than "&" document member of the same patent family the priority date claimed Date of mailing of the international search report Date of the actual completion of the international search 05 -08- 1999 4 August 1999 Authorized officer Name and mailing address of the ISA Swedish Patent Office Hampus Rystedt/CF Box 5055, S-102 42 STOCKHOLM Telephone No. - 46 8 782 25 00 Facsimile No. +46 8 666 02 86

# INTERNATIONAL SEARCH REPORT

Information on patent family members

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